

RESEARCH ARTICLE

Open Access



Nitrous oxide emission by the non-denitrifying, nitrate ammonifier *Bacillus licheniformis*

Yihua Sun¹, Paul De Vos^{1,2} and Kim Heylen^{1*}

Abstract

Background: *Firmicutes* have the capacity to remove excess nitrate from the environment via either denitrification, dissimilatory nitrate reduction to ammonium or both. The recent renewed interest in their nitrogen metabolism has revealed many interesting features, the most striking being their wide variety of dissimilatory nitrate reduction pathways. In the present study, nitrous oxide production from *Bacillus licheniformis*, a ubiquitous Gram-positive, spore-forming species with many industrial applications, is investigated.

Results: *B. licheniformis* has long been considered a denitrifier but physiological experiments on three different strains demonstrated that nitrous oxide is not produced from nitrate in stoichiometric amounts, rather ammonium is the most important end-product, produced during fermentation. Significant strain dependency in end-product ratios, attributed to nitrite and ammonium, and medium dependency in nitrous oxide production were also observed. Genome analyses confirmed the lack of a nitrite reductase to nitric oxide, the key enzyme of denitrification. Based on the gene inventory and building on knowledge from other non-denitrifying nitrous oxide emitters, hypothetical pathways for nitrous oxide production, involving NarG, NirB, qNor and Hmp, are proposed. In addition, all publically available genomes of *B. licheniformis* demonstrated similar gene inventories, with specific duplications of the *nar* operon, *narK* and *hmp* genes as well as NarG phylogeny supporting the evolutionary separation of previously described distinct BAL1 and BAL2 lineages.

Conclusions: Using physiological and genomic data we have demonstrated that the common soil bacterium *B. licheniformis* does not denitrify but is capable of fermentative dissimilatory nitrate/nitrite reduction to ammonium (DNRA) with concomitant production of N₂O. Considering its ubiquitous nature and non-fastidious growth in the lab, *B. licheniformis* is a suitable candidate for further exploration of the actual mechanism of N₂O production in DNRA bacteria and its relevance *in situ*.

Keywords: Dissimilatory nitrate/nitrite reduction to ammonium (DNRA), Fermentation, Nitrate respiration, Denitrification, Ammonification, Nitrite detoxification

Background

Denitrification and dissimilatory nitrate/nitrite reduction to ammonium (DNRA) are two key processes, performed by a wide range of *Bacteria* and *Archaea* as well as some *Eukaryotes* [1], responsible for removal of excess nitrate from the environment. Denitrification is the modular step-wise reduction of fixed nitrogen, nitrate or nitrite to a gaseous form, either nitric oxide (NO),

nitrous oxide (N₂O) and/or dinitrogen gas (N₂). DNRA retains nitrogen in the environment, although N₂O, contributor to both climate change and ozone depletion in the stratosphere, can also be produced as side product. Comprehensive understanding of the identities and activities of microorganisms as well as cellular mechanisms involved in nitrate removal are crucial for improving models that predict fluxes of nitrate, nitrite and N₂O [2]. Although several *Firmicutes* have been known for a long time to be nitrate reducers and N₂O emitters [3–8], their ecological relevance has been minimized over the past two decades based on molecular community surveys

* Correspondence: Kim.Heylen@ugent.be

¹Department of Biochemistry and Microbiology, Laboratory of Microbiology, (LM-Ugent), University of Ghent, K.L. Ledeganckstraat 35, 9000 Ghent, Belgium
Full list of author information is available at the end of the article



using primers not targeting their divergent denitrification [9–12] or DNRA genes [13] (note that recent primers for DNRA do indeed target *Firmicute* genes [14]). Nevertheless, *Firmicutes* and specifically *Bacillus* can be dominant in ecosystems with important nitrate removal activities such as soil [15], animal manure compost [16] and advanced wastewater treatments [17].

Renewed interest in *Bacillus* has revealed many interesting features like (i) the widespread occurrence of nitrate reduction and denitrification in the genus [18], (ii) the gene inventory for both denitrification and DNRA in one microorganism [19, 20], (iii) a novel type of copper-A-dependent, electrogenic nitric oxide reductase (Cu_ANor) [21–24], or (iv) membrane-bound denitrification [25] with a novel organization for the periplasmic nitrate reductase [19, 26]. The most striking observation however is the wide variety of dissimilatory nitrate reduction pathways in members of this genus. The model organism *Bacillus subtilis* uses the cytoplasmic nitrate reductase NarGHI and nitrite reductase NirBD to anaerobically reduce nitrate to ammonium [27, 28], while *Bacillus selenitireducens* produces ammonium via the periplasmic nitrite reductase NrfA [14, 29]. *Bacillus vireti* can do the same but with concomitant N_2O production via Cu_ANor that can be converted to the harmless N_2 with a NosZ-type reductase [30]. On the other hand, *Bacillus azotoformans* and *Bacillus bataviensis* are canonical denitrifiers, the latter lacking the final reductase, but both also encode the NrfA nitrite reductase, making them potential ammonium producers [19]. In addition, these two organisms demonstrate an unusual high level of gene redundancy, i.e. multiple genes or gene copies encoding the same function (*B. azotoformans* encodes three nitrate, two nitrite, four NO and three N_2O reductases) [19]. Considering the modularity of denitrification and DNRA, a multitude of enzyme combinations for nitrate reduction are imaginable, even within one microorganism.

Bacillus licheniformis, a close relative of *B. subtilis*, is widely distributed as a saprophytic organism in the environment, has numerous commercial and agricultural uses (e.g. production of peptide antibiotics, chemicals and proteases, mitigation of fungal pathogens) and some strains, with abortifacient potential or toxin production, might pose a threat to public health. Certain *B. licheniformis* isolates have been described as denitrifiers [5, 18, 28], mostly based on their ability to produce gas from nitrate anaerobically. Many genomes from *B. licheniformis* have been sequenced and described to date [31–36]. However, their lack of genes encoding either a copper- or cytochrome cd_1 -dependent nitrite reductase (NirK or NirS respectively), the key enzyme of denitrification, has gone unnoticed, probably because of limited interest in their anaerobic nitrogen metabolism. We have sequenced and analyzed the genomes of three *B. licheniformis* strains

previously reported to produce N_2O [18], and confirmed the lack of *nirS* or *nirK* in their genomes. In addition, physiological data was gathered demonstrating that *B. licheniformis* does not denitrify but is capable of fermentative dissimilatory nitrate/nitrite reduction to ammonium with concomitant production of N_2O . Both types of data were combined to propose hypothetical pathways for N_2O production, which present new alternative routes for nitrate reduction and N_2O production in members of the genus *Bacillus*.

Methods

Strains and DNA extraction

B. licheniformis LMG 6934, LMG 7559 and LMG 17339 were obtained from the BCCM/LMG bacteria collection. Strains were grown aerobically in trypticase soy broth (TSB) at 37 °C. Cells were harvested after overnight growth and DNA was extracted by the method of Pitcher et al. [37], slightly modified as described previously [38].

Genome sequencing & annotation

Library preparation and genome sequencing was performed by Baseclear B.V. For sequencing, paired-end strategy on the Illumina Genome Analyzer IIx was used yielding average read lengths of 75 bp for LMG 7759 and LMG 17339 and 50 bp for LMG 6934. Automatic trimming (based on a threshold of $Q = 20$ and maximum 2 ambiguous bases) and assembly was performed using CLC Genomics Workbench 6.5. The k-mer and bubble size parameters were varied to maximize the N50 and the number of contig of the resulting assembly for each genome. For the consensus sequence, conflicts were resolved by using quality scores and insertion of ambiguity codes. Functional annotation and metabolic reconstruction was performed with the Rapid Annotation Subsystem Technology (RAST) server [39, 40], using RAST gene calling and allowing frame shift correction, backfilling of gaps and automatic fixing of errors. Assigned functions were checked with pBLAST [41] and InterProScan [42]. Missing genes were searched for in the genome with PSI-BLAST using homologous amino acid sequences. The average nucleotide identity (ANI) was calculated with the ANI calculator (<http://enve-omics.ce.gatech.edu/ani/>) [43].

Growth experiments

Anaerobic growth experiments were performed in TSB and mineral medium (MM), amended with 10 mM potassium nitrate as electron acceptor. Mineral medium was as described by Stanier et al. [44], including 10 mM phosphate buffer, 2.3 mM $(\text{NH}_4)_2\text{SO}_4$, 0.4 mM $\text{MgSO}_4 \cdot 7\text{H}_2\text{O}$, 0.04 mM $\text{CaCl}_2 \cdot 2\text{H}_2\text{O}$, 2.3 mM $(\text{NH}_4)_2\text{SO}_4$, 27 μM EDTA, 25 μM $\text{FeSO}_4 \cdot 7\text{H}_2\text{O}$, 10 μM $\text{ZnSO}_4 \cdot 7\text{H}_2\text{O}$, 25 μM

MnSO₄•H₂O, 3.8 μM CuSO₄•5H₂O, 2 μM Co(N₂O₃)₂•6H₂O, 0.196 μM (NH₄)₆Mo₇O₂₄•24H₂O, supplemented with 30 mM glucose as electron donor. Serum vials (120 ml) were rinsed with 1 M HCl overnight to remove growth inhibiting substances, and subsequently washed four times and rinsed with distilled water before use. Serum vials with 50 ml medium were sealed with black butyl-rubber stoppers. After autoclaving, the headspace of the serum vials was replaced via five cycles of evacuating and refilling with helium. Serum vials were inoculated (1 % v/v) with a suspension of optical density OD₆₀₀ of 1.0 ± 0.05. Each growth experiment was performed in triplicate and non-inoculated media in duplicate were included to check for potential nitrosation reactions in sterile medium. After inoculation, serum vials were incubated at 37 °C, 150 rpm, for 72 h. Preliminary end-point analyses demonstrated that all three strains did not produce N₂ (later confirmed by absence of *nosZ* gene from the genomes, see further) so their dissimilatory nitrate reduction metabolism was explored without addition of acetylene to the headspace. Statistical differences in growth rate and yield of LMG 6934 between TSB and TSB amended with 10 mM nitrate were assessed using the independent t-test after Levene's test for equality of variances, and main and interaction effect of medium and strain on end-product concentration using factorial ANOVA and Least Significant Difference post-hoc testing in SPSS23.

Analytical procedures

Samples of 1 ml were taken from cultures through the rubber septum of serum vials with sterile syringes for growth determination and colorimetric determination of ammonium, nitrate and nitrite. Growth was determined by measuring the optical density OD₆₀₀ of 100 μl sample in duplicate in microtiter plates and standardized to 1 cm pathlength using PathCheck Sensor of the spectrophotometer (Molecular Devices, Spectramax plus 384, USA). For colorimetrics, 500 μl from remaining sample was pretreated with 2.5 ml of 2 M potassium chloride by shaking 1 h at 150 rpm and subsequent filtration (0.2 μm) to extract inorganic nitrogen and remove interfering compounds. Filtered samples were centrifuged at 13000 rpm for 2 min to remove the cells and kept frozen at -20 °C until colorimetric determination. Ammonium concentration was determined with the salicylate-nitroprussidine method (absorption at a wavelength of 650 nm) [45], nitrite and nitrate concentrations were determined with Griess reaction [46] and Griess reaction with cadmium [47, 48] respectively. For end-point measurements, ammonium production was corrected per strain for the amount of ammonium assimilated based on OD₆₀₀ values obtained. Standard curves covered ranges suitable for the tested media and were strictly linear with an R₂ of 0.99.

For determination of N₂O, 1 ml sample of the headspace of serum vials was taken with sterile syringes, and was injected into the gas chromatograph (Compact GC with EZChrom Elite Software, Interscience, Netherlands, 2012). N₂O concentrations were corrected for pressure and solubility based on Henry's law.

Accession numbers

The Whole Genome Shotgun projects of *B. licheniformis* LMG 6934, LMG 7559 and LMG 17339 have been deposited at DDBJ/EMBL/GenBank under the accession numbers AZSY000000000, AZSX000000000, and AZSZ000000000 respectively. The versions described in this paper are the first versions.

Results

Dissimilatory nitrate reduction metabolism

Three genotypically distinct *B. licheniformis* strains (Coorevits, A. & De Vos, P., personal communication) from various origins were selected for determination of their dissimilatory nitrate reduction metabolism based on a previous study that demonstrated their capacity to produce N₂O [18]. *B. licheniformis* LMG 6934 was originally isolated from garden soil, LMG 7559 from flour and LMG 17339 from silage.

Growth of LMG 6934 (Fig. 1a) under anaerobic headspace in TSB amended without and with nitrate (11 mM) commenced after a short lag phase of approximately 3 h, a steep exponential phase followed, with maximal growth achieved after 8.5 h, after which cells sporulated very quickly without a stationary phase (Fig. 1a). Between 6.5 and 7.5 h, growth slowed down probably due to a depletion of the preferential carbon source in the medium with a shift to another electron donor, as this was observed both for fermentative and respiratory growth. The anaerobic growth rates were comparable with and without nitrate as electron acceptor ($\mu_{\text{nitrate}}/\text{h} = 0.189 \pm 0.004 \text{ h}^{-1}$ and $\mu_{\text{ferm}} = 0.179 \pm 0.011 \text{ h}^{-1}$ ($p = 0.264$)), but with significantly different maximal growth yield (OD₆₀₀ of 1.05 ± 0.02 with nitrate and 0.75 ± 0.06 without nitrate ($p = 0.002$)) reflecting the different ATP yield of a respiratory and a fermentative life style. In the presence of nitrate (Fig. 1b), growth was initially supported by nitrate reduction ($\mu_1 = 0.19 \pm 0.004 \text{ h}^{-1}$), with concomitant nitrite production. All nitrate was converted to nitrite but the maximal nitrite peak was probably missed between 5 and 6.5 h of incubation, which was deduced from the onset of N₂O production (0.2 mM N-N₂O at 6.5 h). When nitrate was almost depleted (at 6.5 h, 0.62 mM residual nitrate), nitrite was reduced ($\mu_2 = 0.16 \pm 0.02 \text{ h}^{-1}$), which continued after maximal growth at 8.5 h was achieved and sporulation had started (μ_3). This suggested that nitrite reduction did not support growth during the μ_2 phase, but rather fermentation was responsible for growth after nitrate depletion. During nitrite reduction, a

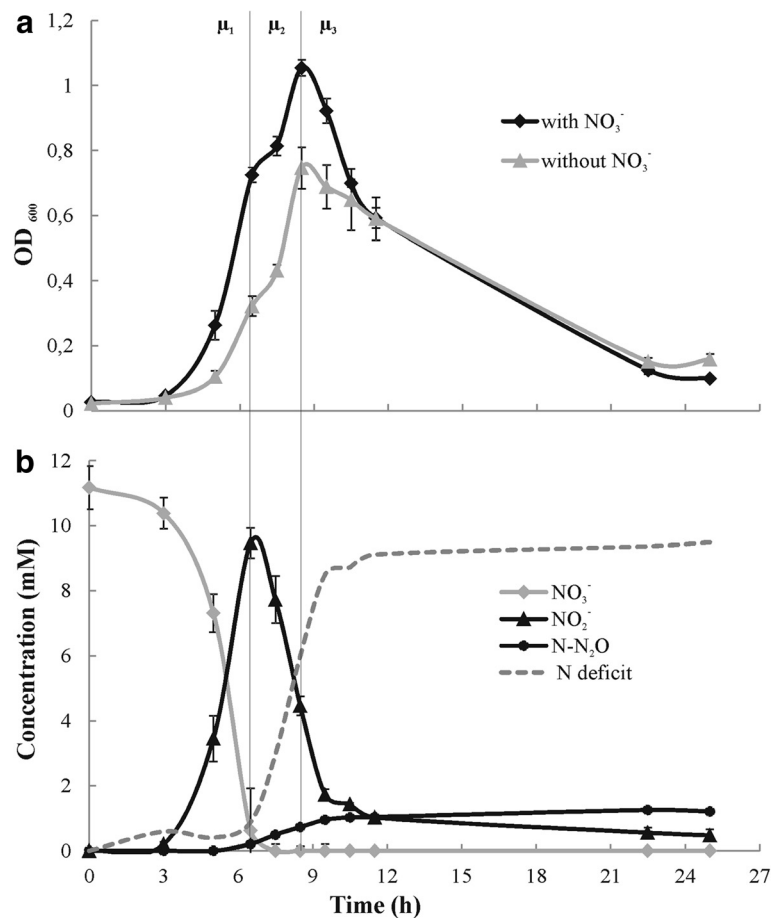


Fig. 1 Anaerobic growth (OD₆₀₀) of *B. licheniformis* LMG 6934 in TSB (a) and nitrate, nitrite and N_2O concentrations (mM)(b) over time. Error bars show standard deviation ($n = 3$). Different growth phases based on primary metabolism in TSB amended with nitrate are marked: μ_1 , respiratory growth with nitrate as electron acceptor; μ_2 , fermentative growth after nitrate is depleted; μ_3 , sporulation. Dashed curve visualizes N deficit caused by the lack of ammonium data

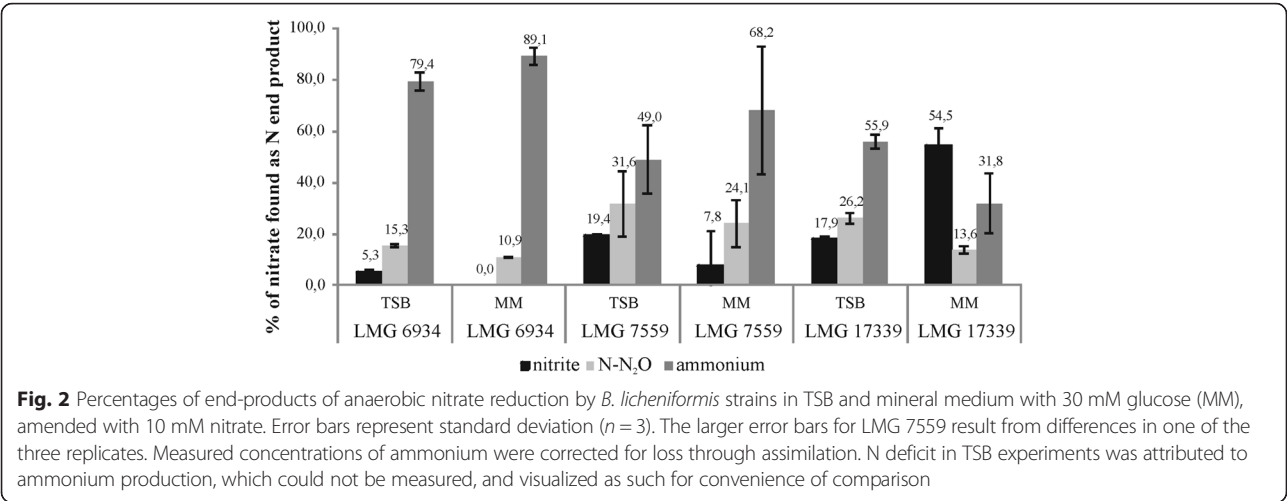
continuous increase in N_2O was observed, with a maximum of 1.3 ± 0.07 mM $\text{N-N}_2\text{O}$ at the end of the incubation (accounting for 12 % of all reduced nitrite), and 0.5 ± 0.18 mM of nitrite remaining in the medium (Fig. 1b). Due to technical constraints and interference of amines from degradation of proteins in the TSB during bacterial growth, ammonium was not monitored during these growth experiments.

To compare the dissimilatory nitrate reduction metabolism of LMG 6934 with those of LMG 7559 and LMG 17339 and to confirm ammonium production from nitrite, end-point experiments after a 72 h-incubation in anaerobic conditions were performed in TSB and mineral medium with 30 mM glucose, both amended with nitrate. Maximal growth of LMG 7559 and LMG 17339 was achieved within 11 h, again immediately followed by a rapid sporulation (data not shown). In mineral media with glucose, most nitrate was converted to ammonium (31.8 to 89.1 %; Fig. 2), confirming the ammonium-producing capacity of all three strains (for TSB, the nitrogen deficit

was attributed to ammonium production, which could not be measured). A significant strain effect on the ratios of end-products was observed ($p \leq 0.008$) (Fig. 2), which after decomposition appeared to be mostly attributed to differences in ammonium and nitrite concentrations. In addition, the amount of $\text{N-N}_2\text{O}$ produced from nitrate was substantially lower in mineral medium than in TSB for all strains (15.8–32.9 % for TSB vs 10.9–24.1 % in MM) ($p = 0.023$). Only for LMG 17339, the medium also had a significant effect on ammonium and nitrite ($p \leq 0.024$). Both strain and medium effect were quite unexpected, as the three strains are closely related and the major carbon source in TSB, namely glucose, is also used in the mineral medium.

Genome analyses

Draft genomes of the three *B. licheniformis* strains were obtained, the genome statistics are given in Table 1. The gene inventory for assimilatory and dissimilatory nitrate reduction and related transport and regulation was



almost identical for the three genomes (Table 2). Details will be given for LMG 6934 and differences with LMG 7559 and LMG 17339 will be highlighted.

The genome of LMG 6934 contained two copies of the *nar* operon (*narGHJ*) coding for the cytoplasmic, membrane-bound nitrate reductase (Table 2; Additional file 1: Figure S1). The two NarG sequences are quite divergent, only sharing 53.7 % amino acid sequence identity. The *nar1* operon (Fig. 3) is located in a gene cluster with downstream the genes for the anaerobic regulatory protein Fnr (Fumarate-Nitrate reductase Regulation) (*fnr1*), a NarK2-type low-affinity nitrate/nitrite antiporter (*narK1*) (Additional file 1: Figure S2), a second Fnr (*fnr2*), a quinol-dependent NO reductase (*qnorB*) and a NO-dependent regulator (*dnrN*). The *nar2* operon (Fig. 3) is immediately downstream of CDS for a second NarK2-type nitrate/nitrite antiporter (*narK2*) (Additional file 1: Figure S2) and a flavohemoprotein (*hmp1*). The two-component nitrate/nitrite sensor regulator system (*narXL*) is encoded downstream of the genes for a nitrite-sensitive transcriptional repressor of NO stress response (*nsrR*) and a second flavohemoprotein (*hmp2*), while genes for a third NarK2-type

nitrate/nitrite antiporter (*narK3*), a formate/nitrite transporter (*nirC*), a second *narL* copy NO reductase activation proteins (*norDQ*), and the global nitrogen regulator (*tnrA*) are found separate on the genome. The gene for NO synthase (*nos*) was also found. Two genes encoding a putative NorV, a flavorubredoxin that could be capable of detoxification of NO to N₂O [49, 50], were also found. However, no gene for the associated oxidoreductase NorW or regulator NorR were found down- and upstream respectively, suggesting that NorV is unlikely to be functional as NO reductase. Nevertheless, all features for nitrate sensing, transport, reduction to nitrite and its regulation are found, as well as for NO reduction to N₂O. In addition, related to nitrogen assimilation, the operon for assimilatory nitrate and nitrite reduction and two genes for AmtB-type ammonium transporters with each upstream the regulatory gene *glnK* are found. Notably, genes for a NirS- or NirK-type nitrite reductase to NO, a NosZ-type N₂O reductase, or a Nrf-type nitrite reductase to ammonium are absent from the genome. The gene inventory and organization for LMG 7559 was identical to LMG 6934 (Table 2). Note that strain LMG 7559 is equivalent to ATTC 9945, for which a complete genome sequence has already been published since the start of our genome analyses [33]. For clarity, both genomes will be included in the remainder of the genome analyses. The genome of LMG 17339 only contained one *nar* operon, two NarK2-type nitrate/nitrite antiporters and one copy of *hmp* and *narL* (associated with *narX*), but for the remainder was identical in gene content and organization to LMG 6934 (Table 2).

Whole genome clustering based on the peptidome content [51, 52], in which the amino acid sequences of a genome are converted to tryptic peptides, i.e. the tryptic peptidome, of all publically available *B. licheniformis* genomes (dd June 2014) confirmed the two generally accepted distinct lineages within *B. licheniformis*, BALI1 and BALI2 [53] (Additional file 1: Figure S3). Average

Table 1 Genome characteristics of three analyzed *B. licheniformis* genomes. Number of coding sequences is based on annotation obtained via RAST

	LMG 6934	LMG 7559	LMG 17339
# contigs (# bp)	53 (4,138,686 bp)	69 (4,341,862 bp)	80 (4,333,151 bp)
N ₅₀	654,545	123,311	102,733
av. read coverage	87.4	82.3	233.7
% G + C	45.9	45.8	46.1
RNA	1 rRNA operon30 tRNA	1 rRNA operon30 tRNA	1 rRNA operon36 tRNA
# coding sequences	4576	4559	4425
accession number	AZSY00000000	AZSX00000000	AZSZ00000000

Table 2 Overview of gene inventory involved in nitrogen metabolism, transport and regulation of *Bacillus licheniformis*

function	protein	gene	locus_tag (gene coordinates)		
			LMG 6934	LMG 7559	LMG 17339
dissimilatory nitrate reduction to nitrite	respiratory nitrate reductase subunit alpha	<i>narG1</i>	LI6934_10240 (contig17_44191_40505)	LI7559_10220 (contig17_109260_112943)	LI17339_03250 (contig01_602796_599110)
		<i>narG2</i>	LI6934_11815 (contig19_3766_7449)	LI7559_18375 (contig29_333921_330235)	-
	respiratory nitrate reductase subunit beta	<i>narH1</i>	LI6934_10235 (contig17_40515_39046)	LI7559_10225 (contig17_112933_114483)	LI17339_03245 (contig01_599120_597651)
		<i>narH2</i>	LI6934_11820 (contig19_7439_8989)	LI7559_18370 (contig29_330245_328776)	-
	respiratory nitrate reductase subunit delta	<i>narJ1</i>	LI6934_10230 (contig17_39028_38486)	LI7559_10230 (contig17_114470_115018)	LI17339_03240 (contig01_597632_597090)
		<i>narJ2</i>	LI6934_11825 (contig19_8976_9524)	LI7559_18365 (contig29_328758_328216)	-
	respiratory nitrate reductase subunit gamma	<i>narI1</i>	LI6934_10225 (contig17_38489_37803)	LI7559_10235 (contig17_115039_115740)	LI17339_03235 (contig01_597093_596407)
		<i>narI2</i>	LI6934_11830 (contig19_9545_10246)	LI7559_18360 (contig29_328219_327533)	-
	assimilatory nitrate reductase large subunit	<i>nasC</i>	LI6934_20135 (contig48_34712_36769)	LI7559_01055 (contig02_88956_91013)	LI17339_17560 (contig10_88671_90728)
		<i>nirB</i>	LI6934_20140 (contig48_36883_39303)	LI7559_01060 (contig02_91127_93547)	LI17339_17565 (contig10_90842_93262)
reduction to ammonium	assimilatory nitrite reductase [NAD(P)H] large subunit	<i>nirD</i>	LI6934_20145 (contig48_39334_39654)	LI7559_01065 (contig02_93578_93898)	LI17339_17560 (contig10_93293_93613)
	assimilatory nitrite reductase [NAD(P)H] small subunit				
transporters	ammonium transport	<i>amt1</i>	LI6934_11075 (contig18_87147_88526)	LI7559_21025 (contig40_60895_59966)	LI17339_05915 (contig03_429269_428058)
		<i>amt2</i>	LI6934_06945 (contig13_69958_71169)	LI7559_12245 (contig20_86027_87406)	LI17339_15165 (contig06_266544_265900)
	nitrate/nitrite transporter (NarK2-type)	<i>nark1</i>	LI6934_10275 (contig17_50582_49398)	LI7559_10215 (contig17_107711_109207)	LI17339_03285 (contig01_609186_608002)
		<i>nark2</i>	LI6934_11810 (contig19_2217_3713)	LI7559_18410 (contig29_340309_339125)	LI17339_16960 (contig09_16054_161849)
		<i>nark3</i>	LI6934_04585 (contig09_42906_44111)	LI7559_1255 (contig21_43506_44711)	-
	formate/nitrite transporter	<i>nirC</i>	LI6934_08170 (contig13_306187_306975)	LI7559_03215 (contig06_176423_177211)	LI17339_04610 (contig03_178114_177326)
	NO reductase large subunit-like protein	<i>qnorB</i>	LI6934_10290 (contig17_51910_54264)	LI7559_18425 (contig29_341636_343990)	LI17339_03300 (contig01_610516_612870)
	NO reductase activation protein	<i>norD1</i>	LI6934_02700 (contig04_119995_118082)	LI7559_00105 (contig01_24308_22395)	LI17339_13375 (contig04_735870_736760)
		<i>norD2</i>	LI6934_02705 (contig04_120896_120006)	LI7559_00110 (contig01_25209_24319)	LI17339_13380 (contig04_736770_738683)
detoxification	flavohemoglobin	<i>hmp1</i>	pLI6934_11805 (contig19_1907_689)	LI7559_10210 (contig17_107401_106181)	LI17339_14225 (contig06_89281_88067)
		<i>hmp2</i>	LI6934_03825 (contig07_12629_11415)	LI7559_14100 (contig23_84124_85338)	-
	NO synthase	<i>nos</i>	LI6934_16440 (contig34_14900_15997)	LI7559_14725 (contig27_17226_16129)	LI17339_20905 (contig20_78781_79878)
regulation	P _{II} -type signal transduction protein	<i>glnK1</i>	LI6934_11070 (contig18_86749_87090)	LI7559_12240 (contig20_85629_85970)	LI17339_05910 (contig03_428039_427689)
		<i>glnK2</i>	LI6934_06950 (contig13_71188_71538)	LI7559_21020 (contig40_59850_59500)	LI17339_15170 (contig06_267014_266677)
	global nitrogen regulator	<i>tnrA</i>	LI6934_15560 (contig31_18775_18443)	LI7559_22170 (contig47_44732_45064)	LI17339_00475 (contig01_88311_87979)

Table 2 Overview of gene inventory involved in nitrogen metabolism, transport and regulation of *Bacillus licheniformis* (Continued)

NO-dependent regulator DnrN or NorA	<i>dnrN</i>	LI6934_10295 (contig17_55022_54309)	LI7559_18430 (contig29_344748_344035)	LI17339_03305 (contig01_613628_612915)
Nitrite-sensitive transcriptional repressor of NO stress response	<i>nsrR</i>	LI6934_03830 (contig07_12887_13318)	LI7559_14095 (contig23_83866_83432)	LI17339_14220 (contig06_87809_87378)
Nitrate/nitrite sensor protein	<i>narX1</i>	LI6934_03855 (contig07_16463_15315)	LI7559_14090 (contig23_83082_83402)	LI17339_14195 (contig06_84234_85382)
Nitrate/nitrite response regulator protein	<i>narL1</i>	LI6934_03850 (contig07_15327_14677)	LI7559_02755 (contig06_92319_91687)	LI17339_14200 (contig06_85379_86023)
	<i>narL2</i>	LI6934_07715 (contig_13_222155_221523)	LI7559_14070 (contig23_80295_81443)	-
transcriptional regulator Crp/Fnr	<i>fnr1</i>	LI6934_10270 (contig17_49251_48538)	LI7559_18405 (contig29_338978_338265)	LI17339_03280 (contig01_607856_607143)
	<i>fnr2</i>	LI6934_10285 (contig17_51086_51754)	LI7559_18420 (contig29_340813_341487)	LI17339_03295 (contig01_609691_610359)

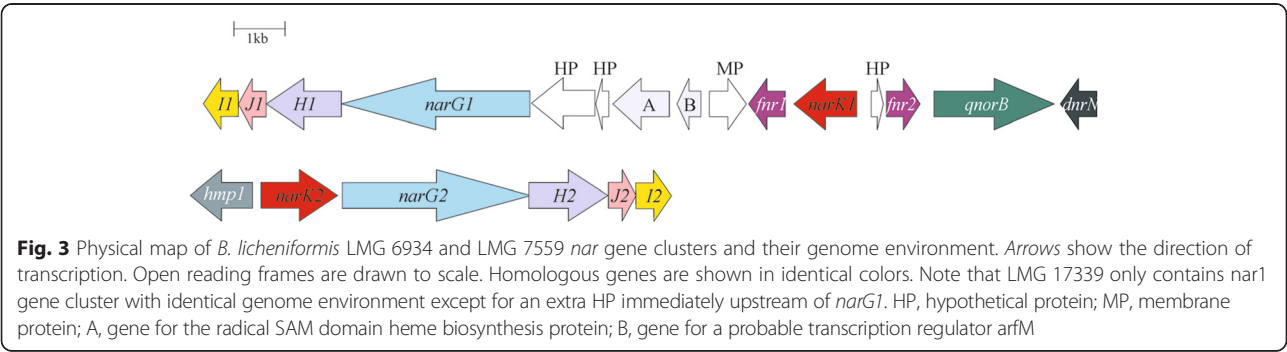
nucleotide identities (ANI) of the genomes from the strains within BALI1 (99.70 % \pm 0.03) and BALI2 (98.92 % \pm 0.03) were well-above the arbitrary 94–95 % cut-off criterion for species delineation, while between group ANI values were ambiguous (94.24 % \pm 0.07) [54, 55]. Interestingly, comparison of the NarG sequences of the three *B. licheniformis* genomes from this study, all other publically available *B. licheniformis* genomes and representatives of other *Bacillus* species showed two distinct clusters, each supported by high bootstrap values (Additional file 1: Figure S1). NarG1 from LMG 6934, NarG2 from LMG 7559 and NarG from LMG 17339 grouped within the BALI 1 cluster, consisting of sequences derived only from *B. licheniformis* and one *Bacillus* sp. NarG2 from LMG 6934 and NarG1 from LMG 7559 fell within cluster BALI2, which also included *B. bataviensis* and one of the two NarGs from *B. azotoformans*. In addition, all *B. licheniformis* genomes from BALI1 consistently harbored only one copy of the *nar* operon, two copies of *narK* and one copy of *hmp*, in contrast to those from BALI2 with two *nar* operons, three *narK* copies and two *hmp* copies (data not shown).

Discussion

General metabolism

Strains belonging to the species *B. licheniformis* have often been considered as denitrifiers [5, 18], based on

their ability to produce gaseous end-products specifically N₂O from nitrate. However, our analyses demonstrated that *B. licheniformis*, like *B. subtilis* [27, 28], is capable of nitrate respiration and fermentative dissimilatory nitrite reduction to ammonium rather than denitrification. All currently available *B. licheniformis* genomes lack a *nirK*- or *nirS*-type nitrite reductase and growth experiments with three strains confirmed that nitrogen gasses were not produced in stoichiometric amounts. *B. licheniformis* first reduced nitrate to nitrite (Fig. 1, μ 1), using the cytoplasmic NarGHI, to support growth accumulating high levels of nitrite before subsequently switching to fermentation after nitrate got depleted. During fermentation (Fig. 1, μ 2), ammonium was produced from nitrite, probably using the NADP-dependent nitrite reductase NirBD (also called NasDE) that can serve for both assimilation and dissimilation. For *B. subtilis* it is still undetermined if nitrite reduction is coupled to energy production through proton motive force [28]. However, since nitrite reduction and production of N₂O continued after growth had ceased, as described before for other non-denitrifying N₂O producers [30, 56, 57], these N conversion seem unrelated to energy conservation. Nevertheless, during fermentative growth, DNRA can serve as an electron-sink allowing re-oxidation of NADH with the generation of one extra ATP by



substrate level phosphorylation for each acetate produced [58, 59].

Hypothesis for NO and N₂O formation

To our knowledge, NO or N₂O production and associated cellular mechanisms have never been described for the model organism *B. subtilis*. Therefore, we built on the knowledge from other model organisms to deduce plausible hypotheses to explain our observations (Fig. 4). Nitrite conversion to NO in *E. coli* was shown to occur only after nitrate was depleted, in presence of molybdate - the cofactor of NarGHI -, continued in *nirB*-mutants [60] but was absent in *narG*-mutants [61], suggesting that the NarGHI had a double function and converted nitrite to NO. Also in *Salmonella enterica* serovar *Typhimurium*, NarGHI was unequivocally responsible for NO generation from nitrite, which was completely eliminated in a *narGHI* mutant [62]. Later mutagenesis experiments in *E. coli* could not confirm the involvement of the cytoplasmic nitrate reductase in NO evolution, probably because the experiments were conducted in the absence of nitrate and thus lacked nitrite formed from NarGHI activity during growth [63]. Rather NirB and NrfA, besides their primary role converting nitrite to ammonium, appeared to be involved in NO production [63], with their relative importance dependent on the nitrite concentration [64, 65]. But, in contrast to *B. vireti* capable of DNRA in combination with NosZ-mediated N₂O reduction [30] but similar to *B. subtilis*, *B. licheniformis* lacks a *nrfA* gene. Furthermore, the activity of NO synthase (NOS), which produces NO from arginine as a defense mechanism against oxidative stress under aerobic conditions in *B. subtilis* [66], is unlikely under our hypoxic test conditions, making the involvement of NarGHI or NirB in the generation of NO from nitrite most plausible (Fig. 4).

Next, NO homeostasis is crucial to limit the toxicity of NO, which is a reactive nitrogen species capable of damaging bacterial proteins, lipids and DNA, and binding to metal centers. NO can freely diffuse through the membrane and can be converted to N₂O in the periplasmic-like space by the quinol-dependent NO reductase qNor (Fig. 4). This reductase is known to be present both in denitrifiers and non-denitrifiers [67–70], including pathogenic bacteria where it is part of their defense mechanism against nitrosative stress. In addition, *B. licheniformis* genomes also encode the flavohemoglobin Hmp. Hmp, found both in cytoplasm and periplasm [71], is known to convert NO to nitrate aerobically and to N₂O anaerobically [72, 73]. However, as the latter conversion is at greatly reduced activity [74], it is unsure whether this enzyme is relevant for NO detoxification in *B. licheniformis* with qNorB; indeed *hmp* appeared not to upregulated in anaerobic conditions at high levels of nitrite in *B. vireti* that contained a copper-dependent NO reductase type 1 [30]. The periplasmic NrfA [75, 76] and the cytoplasmic flavorubredoxin NorV and its associated oxidoreductase NorW [49, 50] that can both anaerobically reduce NO to ammonium and/or N₂O, are not found in *B. licheniformis*. Notably, the gene inventory for anaerobic nitrate and nitrite metabolism in *B. licheniformis* and *B. subtilis* only seems to differ in the presence of a *qnorB* gene in the former organism, making it likely that *B. subtilis* is capable of NO production. This was hinted at by micromolar range N₂O production by *B. subtilis* 1A01 [77] for which the genome is unfortunately not available. Mutagenic and transcriptomic studies are necessary to confirm our hypothetical pathways for NO and N₂O production in *B. licheniformis*.

N end-products: environmental significance and microdiversity

Non-denitrifying nitrate reducers, mostly belonging to *Enterobacteriaceae* or *Bacillaceae*, have been reported to

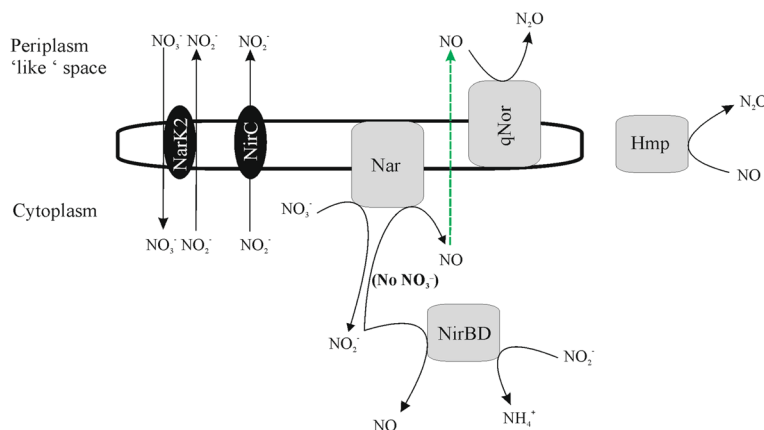


Fig. 4 Proposed hypothetical pathways for anaerobic nitrogen reduction in *B. licheniformis*. Schematic representation of enzymes are given in grey, transporters in black. Diffusion of gaseous NO through cytoplasmic membrane is indicated by the dashed arrow. Soluble Hmp can be located in the cytoplasm and periplasm

reduce typically about 5 to 10 % of nitrate to N_2O , with sometimes high quantities up to 35 %, which evolved mostly after growth has ceased [30, 56, 57, 65, 77]. For *B. licheniformis*, measured N_2O production from nitrate was within these ranges but was nevertheless quite substantial, with up to one-third of all nitrate converted to N_2O (Fig. 2). How environmentally relevant these N_2O emissions from non-denitrifiers are remains difficult to establish without ways to differentiate them from denitrification; in isotope pairing experiments, non-denitrifiers will also produce $^{30}N-N_2O$. In addition, even with mechanistic understanding, deducing specific target genes for molecular surveys will be nearly impossible, as different mechanisms have already been described for a single microorganisms like *E. coli* [78], *S. typhimurium* [62] and *B. vireti* [30] and the genes involved have dual functions, e.g. NarG, NirB and NrfA.

Despite high N_2O emission, most nitrite was indeed converted to ammonium (12.2 – 51.0 %), which is in agreement with previous observations for pure cultures under nitrate limitation (valid for both growth conditions applied here as growth continued via fermentation after nitrate depletion) [30, 56, 77]. Interestingly, the ratio of end-products from nitrate varied quite substantially between all three *B. licheniformis* strains. However, as differences were also apparent between LMG 6934 and LMG 7559, this phenotypic heterogeneity could not be linked the specific gene duplications in BALI2 genomes. An alternative explanation might be distinct regulatory motifs in the promotor regions of the genes involved, although the same regulatory genes were encoded in the three genomes with identical relative genome locations (Table 2). It is long been accepted that closely related bacteria do not necessarily share the capacity to denitrify, and even when they do, can have different denitrifying phenotypes. However, our data suggest that phenotypic heterogeneity or niche differentiation between closely related strains, which has recently been reported for N_2O production in *Bacillus* [79], *Thauera* [80] and *Methylobacillus* [81], might not always be linked to genetic variation.

Conclusions

Using physiological and genomic data we have demonstrated that the common soil bacterium *B. licheniformis* does not denitrify but is capable of fermentative dissimilatory nitrate/nitrite reduction to ammonium with concomitant production of N_2O . Based on the genomic inventory, alternative routes for N_2O production, similar to those in *Enterobacteriaceae* and thus far unreported in bacilli, were proposed. Significant strain-dependent differences were found between three closely related strains that could not be linked to genetic features. Considering its ubiquitous nature and non-fastidious growth in the lab, *B. licheniformis* is a suitable candidate for further

exploration of the uncertainty of the mechanism of N_2O production in DNRA bacteria and its relevance *in situ*.

Availability of supporting data

All the supporting data are included as additional files.

Additional file

Additional file 1: Figure S1. Phylogenetic tree of full-length NarG. Sequences were taken from genomes included in the manuscript as well as all from publically available *B. licheniformis* genomes (dd June 2014) and genomes from other *Bacillus* species, protein ID or locus tag is given between brackets. The evolutionary history was inferred using the Neighbor-Joining method. The optimal tree with the sum of branch length = 1,68972248 is shown. The percentage of replicate trees in which the associated taxa clustered together in the bootstrap test (500 replicates) are shown next to the branches. The evolutionary distances were computed using the Poisson correction method and are in the units of the number of amino acid substitutions per site. All positions containing gaps and missing data were eliminated. There were a total of 1094 positions in the final dataset. Evolutionary analyses were conducted in MEGA6.

Figure S2. Phylogenetic tree of full-length NarK. Sequences were taken from several *Bacillus* species, as well as reference genomes, locus tag is given between brackets. The evolutionary history was inferred using the Neighbor-Joining method. The optimal tree with the sum of branch length = 6,03046656 is shown. The percentage of replicate trees in which the associated taxa clustered together in the bootstrap test (500 replicates) are shown next to the branches. The tree is drawn to scale, with branch lengths in the same units as those of the evolutionary distances used to infer the phylogenetic tree. The evolutionary distances were computed using the Poisson correction method and are in the units of the number of amino acid substitutions per site. All positions containing gaps and missing data were eliminated. There were a total of 358 positions in the final dataset. Evolutionary analyses were conducted in MEGA6.

Figure S3. Whole genome clustering based on a similarity matrix using the peptidome content, in which amino acid sequences of a genome are converted to tryptic peptides, i.e. the tryptic peptidome. All publically available *B. licheniformis* genomes (dd June 2014) were included and are designated by their strain number: 9945A (accession number NC_021362), ATCC 14580 (accession number NC_006270), DSM 13 (=ATCC 14580; accession number NC_006322), 5-2-D (accession number NZ_AJLW01000000), F1-1 (accession number NZ_AZSL01000000), 3F-3 (accession number NZ_JFYM01000000), F2-1 (accession number NZ_AZSM01000000), WX-02 (accession number NZ_JH636050), 10-1-A (accession number NZ_AJLV01000001), CGMCC 3963 (accession number NZ_AMWQ01000000), CG-B52 (accession number NZ_AVEZ01000000). Analyses were performed using the Peptidome tool (<http://unipect.ugent.be/>). (DOCX 29 kb)

Abbreviations

DNRA: Dissimilatory nitrate/nitrite reduction to ammonium; TSB: Trypticase soy broth; RAST: Rapid annotation subsystem technology; ANI: Average nucleotide identity; MM: Mineral medium; OD: Optical density.

Competing interests

The authors declare to have no competing interests.

Authors' contributions

YH performed the growth experiments, data analyses and wrote the paper; KH conceived the study, performed genome data analyses and wrote the paper; PDV helped with interpretation of data and revised the paper. All authors have read and approved the manuscript.

Acknowledgements

YS was funded by Chinese Scholarship Council (File number 201206330054) and BOF CSC co-funding from Ghent University (grant 01SC2713). KH was funded by the Fund for Scientific Research (FWO), Flanders for a position as postdoctoral research fellows (grants FWO11/PDO/0840 and FWO15/PDOH1/

084). Genome sequencing was also funded by FWO (grant FWO11/KAN/043). This research was also supported by Ghent University Research Council (GOA project 01G01911). The funders had no role in study design, data collection and interpretation, or the decision to submit the work for publication.

Author details

¹Department of Biochemistry and Microbiology, Laboratory of Microbiology, (LM-Ugent), University of Ghent, K.L. Ledeganckstraat 35, 9000 Gent, Belgium. ²BCCM/LMG Bacteria Collection, K.L. Ledeganckstraat 35, 9000 Gent, Belgium.

Received: 13 October 2015 Accepted: 6 January 2016

Published online: 19 January 2016

References

- Zumft WG. Cell biology and molecular basis of denitrification. *Microbiol Mol Biol Rev.* 1997;61(4):533–616.
- Canfield DE, Glazer AN, Falkowski PG. The evolution and future of earth's nitrogen cycle. *Science.* 2010;330:192–6.
- Pichinoty F, de Barjac H, Mandel M, Asselineau J. Description of *Bacillus azotoformans* sp. nov. *Int J Syst Bacteriol.* 1983;33:660–2.
- Pichinoty F, Durand M, Job C. Etude morphologique, physiologique et taxonomique de *Bacillus azotoformans*. *Can J Microbiol.* 1978;24:608–17.
- Pichinoty F, Garcia JL, Job C, Durand M. La dénitrification chez *Bacillus licheniformis*. *Can J Microbiol.* 1978;24:45–9.
- Pichinoty F, Mandel M, Garcia JL. The properties of novel mesophilic denitrifying *Bacillus* cultures found in tropical soils. *J Gen Microbiol.* 1979;115:419–30.
- Denariuz G, Payne WJ, Legall J. A halophilic denitrifier, *Bacillus halodenitrificans* sp. nov. *Int J Syst Bacteriol.* 1989;39:145–51.
- Payne WJ. Denitrification. New York: John Wiley & Sons; 1981.
- Braker G, Fesefeldt A, Witzel K-P. Development of PCR primer systems for amplification of nitrite reductase genes (*nirK* and *nirS*) to detect denitrifying bacteria in environmental samples. *Appl Environ Microbiol.* 1998;64(10):3769–75.
- Throback IN, Enwall K, Jarvis A, Hallin S. Reassessing PCR primers targeting *nirS*, *nirK* and *nosZ* genes for community surveys of denitrifying bacteria with DGGE. *FEMS Microbiol Ecol.* 2004;49:401–17.
- Penton CR, Johnson TA, Quensen JF, Iwai S, Cole JR, Tiedje JM. Functional genes to assess nitrogen cycling and aromatic hydrocarbon degradation: primers and processing matter. *Front Microbiol.* 2013;4:279.
- Wei W, Isobe K, Nishizawa T, Zhu L, Shiratori Y, Ohte N, et al. Higher diversity and abundance of denitrifying microorganisms in environments than considered previously. *ISME J.* 2015;9(9):1954–65.
- Mohan SB, Schmid M, Jetten M, Cole J. Detection and widespread distribution of the *nrfA* gene encoding nitrite reduction to ammonia, a short circuit in the biological nitrogen cycle that competes with denitrification. *FEMS Microbiol Ecol.* 2004;49:433–43.
- Welsh A, Chee-Sanford J, Connor L, Löffler F, Sanford R. Refined *NrfA* phylogeny improves PCR-based *nrfA* gene detection. *Appl Environ Microbiol.* 2014;80:2110–9.
- Felske A, Akkermans ADL, De Vos WM. *In Situ* detection of an uncultured predominant *Bacillus* in Dutch Grassland soils. *Appl Environ Microbiol.* 1998;34(11):4588–90.
- Maeda K, Hanajima D, Toyoda S, Yoshida N, Morioka R, Osada T. Microbiology of nitrogen cycle in animal manure compost. *Microb Biotechnol.* 2011;4:700–9.
- Park SJ, Yoon JC, Shin KS, Kim EH, Yim S, Cho YJ, et al. Dominance of endospore-forming bacteria on a rotating activated *Bacillus* contactor biofilm for advanced wastewater treatment. *J Microbiol (Seoul, Korea).* 2007;45(2):113–21.
- Verbaendert I, Boon N, De Vos P, Heylen K. Denitrification is a common feature among members of the genus *Bacillus*. *Syst Appl Microbiol.* 2011;34:385–91.
- Heylen K, Keltjens J. Redundancy and modularity in membrane-associated dissimilatory nitrate reduction in *Bacillus*. *Front Microbiol.* 2012;3:371.
- Nielsen M, Schreiber L, Finster K, Schramm A. Draft genome sequence of *Bacillus azotoformans* MEV2011, a (Co-) denitrifying strain unable to grow with oxygen. *Stand Genomic Sci.* 2014;9:23.
- Lu S, Suharti, Vries S, Moëne-Loccoz P. Two CO molecules can bind concomitantly at the diiron site of NO reductase from *Bacillus azotoformans*. *J Am Chem Soc.* 2004;126:15332–3.
- Suharti, Heering HA, de Vries S. NO reductase from *Bacillus azotoformans* is a bifunctional enzyme accepting electrons from menaquinol and a specific endogenous membrane-bound cytochrome c551. *Biochemistry.* 2004;43:13487–95.
- Suharti, Strampaard MJF, Schröder I, de Vries S. A novel copper A containing menaquinol NO reductase from *Bacillus azotoformans*. *Biochemistry.* 2001;40:2632–9.
- Al-Attar S, de Vries S. An electrogenic nitric oxide reductase. *FEBS Lett.* 2015; 589(16):2050–7.
- Suharti, de Vries S. Membrane-bound denitrification in the Gram-positive bacterium *Bacillus azotoformans*. *Biochem Soc Trans.* 2005;33(1):130–3.
- Simon J, Klotz MG. Diversity and evolution of bioenergetic systems involved in microbial nitrogen compound transformations. *Biochim Biophys Acta.* 2013;1827(2):114–35.
- Nakano MM, Hoffmann T, Zhu Y, Jahn D. Nitrogen and oxygen regulation of *Bacillus subtilis* *nasDEF* encoding NADH-dependent nitrite reductase by TnrA and ResDE. *J Bacteriol.* 1998;180(20):5344–50.
- Nakano MM, Zuber P. Anaerobic growth of a "strict aerobe" (*Bacillus subtilis*). *Annu Rev Microbiol.* 1998;52:165–90.
- Switzer Blum J, Burns Bindi A, Buzzelli J, Stolz JF, Oremland RS. *Bacillus arsenicoselenatis*, sp. nov., and *Bacillus selenitireducens*, sp. nov.: two haloalkaliphiles from Mono Lake, California that respire oxyanions of selenium and arsenic. *Arch Microbiol.* 1998;171(1):19–30.
- Mania D, Heylen K, van Spanning RJ, Frostegard A. The nitrate-ammonifying and *nosZ* carrying bacterium *Bacillus vireti* is a potent source and sink for nitric and nitrous oxides under high nitrate conditions. *Environ Microbiol.* 2014;16:3196–210.
- Dhakal R, Seale RB, Deeth HC, Craven H, Turner MS. Draft genome comparison of representatives of the three dominant genotype groups of dairy *Bacillus licheniformis* strains. *Appl Environ Microbiol.* 2014;80(11):3453–62.
- Li L, Su F, Wang Y, Zhang L, Liu C, Li J, et al. Genome sequences of two thermophilic *Bacillus licheniformis* strains, efficient producers of platform chemical 2,3-butanediol. *J Bacteriol.* 2012;194(15):4133–4.
- Rachinger M, Volland S, Meinhardt F, Daniel R, Liesegang H. First Insights into the Completely Annotated Genome Sequence of *Bacillus licheniformis* Strain 9945A. *Genome Announc.* 2013; 1(4):00525–13.
- Rey MW, Ramaiya P, Nelson BA, Brody-Karpin SD, Zaretsky EJ, Tang M, et al. Complete genome sequence of the industrial bacterium *Bacillus licheniformis* and comparisons with closely related *Bacillus* species. *Genome Biol.* 2004;5:R77.
- Veith B, Herzberg C, Steckel S, Feesche J, Maurer KH, Ehrenreich P, et al. The complete genome sequence of *Bacillus licheniformis* DSM13, an organism with great industrial potential. *J Mol Microbiol Biotechnol.* 2004;7:204–11.
- Wu Q, Peng S, Yu Y, Li Y, Xu Y. Genome Sequence of *Bacillus licheniformis* CGMCC3963, a Stress-Resistant Strain Isolated in a Chinese Traditional Solid-State Liquor-Making Process. *Genome Announcements.* 2013; 1(1):00060–12.
- Pitcher DG, Saunders NA, Owen RJ. Rapid extraction of bacterial genomic DNA with guanidium thiocyanate. *Lett Appl Microbiol.* 1989;8:151–6.
- Heyndrickx M, Vauterin L, Vandamme P, Kersters K, De Vos P. Applicability of combined amplified ribosomal DNA restriction analysis (ARDRA) patterns in bacterial phylogeny and taxonomy. *J Microbiol Meth.* 1996;26:247–59.
- Overbeek R, Olson R, Pusch GD, Olsen GJ, Davis JJ, Disz T, et al. The SEED and the Rapid Annotation of microbial genomes using Subsystems Technology (RAST). *Nucleic Acids Res.* 2014;42(Database issue):D206–214.
- Aziz RK, Bartels D, Best AA, DeJongh M, Disz T, Edwards RA et al. The RAST Server: Rapid Annotations using Subsystems Technology. *BMC Genomics.* 2008; 9(75):doi:10.1186/1471-2164-1189-1175.
- Altschul SF, Madden TL, Schaffer AA, Zhang J, Zhang Z, Miller W, et al. Gapped BLAST and PSI-BLAST: a new generation of protein database search programs. *Nucleic Acids Res.* 1997;25:3389–402.
- Zdobnov EM, Apweiler R. InterProScan - an integration platform for the signature-recognition methods in InterPro. *Bioinformatics.* 2001;17(9):847–8.
- Goris J, Konstantinidis KT, Klappenbach JA, Coenye T, Vandamme P, Tiedje JM. DNA–DNA hybridization values and their relationship to whole-genome sequence similarities. *Int J Syst Evol Microbiol.* 2007;57:81–91.
- Stanier RY, Palleroni NJ, Doudoroff M. The aerobic pseudomonads: a taxonomic study. *J Gen Microbiol.* 1966;43(2):159–271.
- Baethgen WE, Alley MM. A manual colorimetric procedure for measuring ammonium nitrogen in soil and plant Kjeldahl digests. *Commun Soil Sci Plant Anal.* 1989;20:9–10.

46. Griess P. Bemerkungen zu der abhandlung der H.H. Weselsky und Benedikt "Ueber einige azoverbindungen". Chem Ber. 1879;12:426–8.
47. Cataldo DA, Haroon M, Schrader LE, Youngs VL. Rapid colorimetric determination of nitrate in plant-tissue by nitration of salicylic-acid. Commun Soil Sci Plant Anal. 1975;6:71–80.
48. Navarro-Gonzalez JA, Garcia-Benayas C, Arenas J. Semiautomated measurement of nitrate in biological fluids. Clin Chem. 1998;44(3):679–81.
49. Gomes CM, Giuffrè A, Forte E, Vicente JB, Saraiva LM, Brunori M, et al. A novel type of nitric-oxide reductase. *Escherichia coli* flavorubredoxin. J Biol Chem. 2002;277(28):25273–6.
50. van Wonderen JH, Burlat B, Richardson DJ, Cheesman MR, Butt JN. The nitric oxide reductase activity of cytochrome c nitrite reductase from *Escherichia coli*. J Biol Chem. 2008;283:9587–94.
51. Mesuere B, Devreese B, Debyser G, Aerts M, Vandamme P, Dawyndt P. Unipept: tryptic peptide-based biodiversity analysis of metaproteome samples. J Proteome Res. 2012;11(12):5773–80.
52. Mesuere B, Debyser G, Aerts M, Devreese B, Vandamme P, Dawyndt P. The Unipept metaproteomics analysis pipeline. Proteomics. 2015;15(8):1437–42.
53. De Clerck E, De Vos P. Genotypic diversity among *Bacillus licheniformis* strains from various sources. FEMS Microbiol Lett. 2004;231(1):91–8.
54. Konstantinidis KT, Tiedje JM. Genomic insights that advance the species definition for prokaryotes. Proc Natl Acad Sci U S A. 2005;102:2567–72.
55. Konstantinidis KT, Ramette A, Tiedje JM. The bacterial species definition in the genomic era. Phil Trans R Soc B. 2006;361:1929–40.
56. Smith MS, Zimmerman K. Nitrous oxide production by nondenitrifying soil nitrate reducers. Soil Sci Soc Am J. 1981;45(5):865–71.
57. Bleakley BH, Tiedje JM. Nitrous oxide production by organisms other than nitrifiers or denitrifiers. Appl Environ Microbiol. 1982;44(6):1342–8.
58. Cole J. Nitrate reduction to ammonia by enteric bacteria: redundancy, or a strategy for survival during oxygen starvation? FEMS Microbiol Lett. 1996;136(1):1–11.
59. Polcyn W, Podeszwa J. Coordinate induction of dissimilatory ammonification and fermentative pathways in rhizobia. Antonie Van Leeuwenhoek. 2009;96(1):79–87.
60. Smith MS. Dissimilatory reduction of NO_2^- to NH_4^+ and N_2O by a soil *Citrobacter* sp. Appl Environ Microbiol. 1982;43:854–60.
61. Smith MS. Nitrous oxide production by *Escherichia coli* is correlated with nitrate reductase activity. Appl Environ Microbiol. 1983;45(5):1545–7.
62. Gilberthorpe NJ, Poole RK. Nitric oxide homeostasis in *Salmonella typhimurium*: roles of respiratory nitrate reductase and flavohemoglobin. J Biol Chem. 2008;283(17):11146–54.
63. Corker H, Poole RK. Nitric oxide formation by *Escherichia coli*. Dependence on nitrite reductase, the NO-sensing regulator Fnr, and flavohemoglobin Hmp. J Biol Chem. 2003;278(34):31584–92.
64. Wang H, Robert O, Hall J, Gunsalus P. The *nrfA* and *nirB* nitrite reductase operons in *Escherichia coli* are expressed differently in response to nitrate than to nitrite. J Bacteriol. 2000;182(20):5813–22.
65. Rowley G, Hensen D, Felgate H, Arkenberg A, Appia-Ayme C, Prior K, et al. Resolving the contributions of the membrane-bound and periplasmic nitrate reductase systems to nitric oxide and nitrous oxide production in *Salmonella enterica* serovar Typhimurium. Biochem J. 2012;441(2):755–62.
66. Adak S, Aulak KS, Stuehr DJ. Direct evidence for nitric oxide production by a nitric-oxide synthase-like protein from *Bacillus subtilis*. J Biol Chem. 2002;277(18):16167–71.
67. Büsch A, Friedrich B, Cramm R. Characterization of the *norB* gene, encoding nitric oxide reductase, in the nondenitrifying cyanobacterium *Synechocystis* sp. Strain PCC6803. Appl Environ Microbiol. 2002;68(2):668–72.
68. Cramm R, Siddiqui RA, Friedrich B. Two isofunctional nitric oxide reductases in *Alcaligenes eutrophus* H16. J Bacteriol. 1997;197(21):6769–77.
69. Philippot L. Denitrifying genes in bacterial and Archaeal genomes. Biochim Biophys Acta. 2002;1577:355–76.
70. Heylen K, Vanparrys B, Gevers D, Wittebolle L, Boon N, De Vos P. Nitric oxide reductase (*norB*) gene sequence analysis reveals discrepancies with nitrite reductase (*nir*) gene phylogeny in cultivated denitrifiers. Environ Microbiol. 2007;9(4):1072–7.
71. Vasudevan SG, Tang P, Dixon NE, Poole RK. Distribution of the flavohaemoglobin, HMP, between periplasm and cytoplasm in *Escherichia coli*. FEMS Microbiol Lett. 1995;125(2-3):219–24.
72. Gardner PR, Gardner AM, Martin LA, Salzman AL. Nitric oxide dioxygenase: an enzymic function for flavohemoglobin. Proc Natl Acad Sci U S A. 1998;95(18):10378–83.
73. Kim SO, Orii Y, Lloyd D, Hughes MN, Poole RK. Anoxic function for the *Escherichia coli* flavohaemoglobin (Hmp): reversible binding of nitric oxide and reduction to nitrous oxide. FEBS Lett. 1999;445(2-3):389–94.
74. Mills CE, Sedelnikova S, Soballe B, Hughes MN, Poole RK. *Escherichia coli* flavohaemoglobin (Hmp) with equistoichiometric FAD and haem contents has a low affinity for dioxygen in the absence or presence of nitric oxide. Biochem J. 2001;353(Pt 2):207–13.
75. Poock SR, Leach ER, Moir JWB, Cole JA, Richardson DJ. Respiratory detoxification of nitric oxide by the cytochrome c nitrite reductase of *Escherichia coli*. J Biol Chem. 2002;277:23664–9.
76. Costa C, Macedo A, Moura I, Moura JGG, Le Gall J, Berlier Y, et al. Regulation of the hexaheme nitrite/nitric oxide reductase of *Desulfovibrio desulfuricans*, *Wolinella succinogenes* and *Escherichia coli*. FEBS Lett. 1990;276:67–70.
77. Stremiška MA, Felgate H, Rowley G, Richardson DJ, Baggs EM. Nitrous oxide production in soil isolates of nitrate-ammonifying bacteria. Environ Microbiol Rep. 2012;4(1):66–71.
78. Vine CE, Cole JA. Nitrosative stress in *Escherichia coli*: reduction of nitric oxide. Biochem Soc Trans. 2011;39:313–5.
79. Jones CM, Welsh A, Throback IN, Dorsch P, Bakken LR, Hallin S. Phenotypic and genotypic heterogeneity among closely related soil-borne N- and N₂O-producing *Bacillus* isolates harboring the *nosZ* gene. FEMS Microbiol Ecol. 2011;76(3):541–552.
80. Liu B, Mao Y, Bergaust L, Bakken LR, Frostegard A. Strains in the genus *Thauera* exhibit remarkably different denitrification regulatory phenotypes. Environmental microbiology. 2013;15(10):2816–2828.
81. Hoefman S, van der Ha D, Boon N, Vandamme P, De Vos P, Heylen K. Niche differentiation in nitrogen metabolism among methanotrophs within an operational taxonomic unit. BMC microbiology. 2014;14(1):83.

Submit your next manuscript to BioMed Central and we will help you at every step:

- We accept pre-submission inquiries
- Our selector tool helps you to find the most relevant journal
- We provide round the clock customer support
- Convenient online submission
- Thorough peer review
- Inclusion in PubMed and all major indexing services
- Maximum visibility for your research

Submit your manuscript at
www.biomedcentral.com/submit

